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<pre> CCCCCCCCCCCCACCCCTGGACCTGCCTCCCCGGTGCACCCGCACAGCCCTTCCGCGTCAGCAGTCAGTGATCAGGGCCAAAGCCGTCAGTCACCA C S C E P V E D Q A Y C E A D V V I R A K A V S X X E 100 AGTGGACTCTGCAAAACACATTATGGCAACCCCTATCAGAACGATCCAGATATGACATCAAGCAGATAAAAGATGTTCAAGGGCCCTGCAAAACGATATAGAC V D S C N D I Y C G P I K R I Q Y E K Q I K M F X G P E L D I E 200 TTTATCTACAGCCCCCTCTCTCGCCAGCTGCTGCGGCTCTCGACCTGGACAAAACAGCAATATCTCATTCGACAAAAGCCGACGGGGACGGCA V I T T A P E S A V C C V S L D V C C K E T L I A G K A E G D C 300 AGATTCACATCACCCCTCTGACTTCATCTGTCCTCCCTGGCCACCCCTCACACCCACACCCAAACAGAACGCCCTGCAAAACAGCTACCCACATGGCTTCCGACTG K M I T Z L C P W D S L K E Y T Q M H Y T Q M G C E 400 CACATCAACCCCCCTCCCCCATGATCCGGTCTACATCTCTCTCCGGACAGCTGGCTCTGGATGGACTGGCTCACAGAACAGAACATCAACGGGACCCAG K I T Z B C M I P C T S A F D E C L E M D W V T E K H I Z G E Q 500 CCCCAGCTCTCCGCTGCGATCACACACTGACTCTCTCTGGCTGCGTGGACGCTCTGGACGCTGGCTCACAGAACAGAACATCAACGGGACCCAG A K F Y A C T E R S D C S C A V T R G A A D P E K Q E F L D I E D P 600 AACCCAGCCCTCAACCCCGCTGGCTGCGAACCTGCGAAAAAAACCGCTCAACGGCTTCCACCTGCGATCCCTTCTGGCAAAACACCATGATAA 700 ACACTCATCCCCATGGCTCCAAAATTAATATC </pre>					
(57) Abstract					
<p>The present invention is an isolated protein of 21,600 daltons which binds to both latent and activated type IV collagenase with high affinity at 1:1 molar stoichiometry, thereby abolishing enzyme activity. The protein is purified by affinity chromatography on solid phase metalloproteinase, or solid phase metalloproteinase substrates which bind the enzyme-inhibitor complex. The complete primary structure of this protein (initially called CSC-21K), as determined by sequencing overlapping peptides spanning the entire protein, reveals homology with a protein called TIMP, Tissue Inhibitor of Metalloproteinases. In addition, a cDNA for this novel inhibitor, now designated TIMP-2, was cloned from a melanoma cell and its sequence was compared with that of human TIMP-1. Northern blots of melanoma cell mRNA showed two distinct transcripts of 0.9 kb and 3.5 kb which are down-regulated by transforming growth factor-beta, and are unchanged by phorbol ester treatment. The figure 7 shows the cDNA and corresponding protein sequence of TIMP-2.</p>					

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MATRIX METALLOPROTEINASE INHIBITOR PEPTIDES

Field of the Invention

This invention relates to isolated proteins or 5 peptides useful for inhibition of matrix metalloproteinases. Specifically, this invention relates to a novel protein, isolated from conditioned media of cultured human tumor cells, which binds with high affinity to matrix metalloproteinase enzymes and analogs thereof. The natural 10 protein is defined by a novel amino acid sequence including specific positions of cysteine residues. This invention further relates to a novel means of purifying matrix metalloproteinase inhibitors using metalloproteinase affinity chromatography.

15 Background of the Invention

The collagenase family of enzymes are a group of neutral metalloproteinases, also known as matrix metalloproteinases, which are secreted in the zymogen form and degrade both the collagenous and noncollagenous 20 components of the extracellular matrix. All require a metal ion (calcium and/or zinc) for hydrolytic activity, and all are secreted in the latent pre-enzyme form. Members of this collagenase gene family include: the interstitial collagenases, which degrade collagen types I, II and III and 25 have been characterized with respect to substrate specificity and requirements for activation (Stricklin, G.P., Jeffrey, J.J., Rosewit, W.T., and Eisen, A.Z., 1983, Biochemistry 22, 61-68; Goldberg, G.I., Wilhelm, S., Kronberger, A., Bauer, E.A., Grant, G.A., and Eisen, A.Z., 1986, J. Biol. Chem. 261, 30 6600-6605; Hasty, K.A., Jeffrey, J.J., Hibbs, M.S., and Welgus, H.G., 1987, J. Biol. Chem. 262, 10048-1052; Fields, G.B., Van Wart, H.E., and Birkedal-Hansen, H., 1987, J. Biol. Chem. 262, 6221-6226; Grant, G.A., Eisen, A.Z., Marmer, B.L. Rosweit, W.T., and Goldberg, G.I., 1987, J. Biol. Chem. 262, 35 5886-5889); stromelysin, which degrades proteoglycans, glycoproteins, and the non-helical portions of collagenous molecules (Wilhelm, S.M., Collier, I.E., Kronberger, A.,

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Eisen, A.Z., Marmer, B.L., Grant, G.A., Bauer, E., and Goldberg, G.I., 1987, Proc. Natl. Acad. Sci. U.S.A. 84, 6725-6729; Whitman, S.E., Murphy, G., Angel, P., Rahmsdorf, H.-J., SMith, B.J., Lyons, A., Harris, T.J.T., Reynolds. J.J., 5 Herrlich, P. and Docherty, A.J.P., 1986, Biochem. J. 240, 913-916); and type IV collagenase, which degrades pepsin-resistant triple-helical type IV collagen and interstitial collagens (gelatin). Type IV collagenase has been identified in human tumor cells (Liotta, L.A., Kleinerman, J., 10 Catanzaro, P., and Rynbrandt, D., 1977, J. Natl. Cancer Inst. 58, 1427-1439; Turpeenniemi-Hujanen, T., and Tryggvason, K., 1982, Int. J. Cancer 30p, 669-673; Liotta, L.A., Abe, S., Gehron-Robey, P., and Martin, G.R., 1979, Proc. Natl. Acad. Sci. U.S.A. 76 2268-2272; Liotta, L.A., Tryggvasson, K., 15 Garbisa, S., Hart, I., Foltz, C.M., and Shafie, S., 1980, Nature (London) 284, 67-68; Collier, I.E., Wilhelm, S.M., Eisen, A.Z., Marmer, B.L., Grant, G.A., Seltzer, J.L., Kronberger, A., He., C., Bauer, E.A., and Goldberg, G.I., 1988, J. Biol. Chem. 263, 6579-6587), endothelial cells 20 (Kalebic, T., Barbisa, S., Glaser, B., and Liotta, L.A., 1983, Science 221, 281-283), bone (Murphy, G., McAlpine, C.G., Poll, C.T., and Reynolds, J.J., 1985, Biochem. Biophys. Acta 831, 49-58), fibroblasts (Collier, I.E., Wilhelm, S.M., Eisen, A.Z., Marmer, B.L., Grant, G.A., Seltzer, J.L., 25 Kronberger, A., He., C., Bauer, E.A., and Goldberg, G.I., 1988, J. Biol. Chem. 263, 6579-6587), polymorphonuclear leukocytes (Uitto, V.J., Schwartz D., and Veis, A., 1980, Eur. J. Biochem. 105, 409-417) and macrophages (Garbisa, S., Ballin, M., Daga-Giordini, D., Fastelli, G., Naturale, M., 30 Negro, A., Semenzato, G., and Liotta, L.A., 1986, J. Biol. Chem. 261, 2369-2375). This enzyme is a neutral metalloproteinase of 68 to 72 kilodaltons which is secreted in zymogen form (Liotta, L.A., Abe, S., Gehron-Robey, P., and Martin, G.R., 1979, Proc. Natl. Acad. Sci. U.S.A. 76, 2268-35 2272; Liotta, L.A., Tryggvasson, K., Garbisa, S., Gehron-Robey, P., and Abe, S., 1981, Biochemistry 20, 100-104; Salo, T., Liotta, L.A., and Tryggvasson, K., 1983, J. Biol. Chem.

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258, 3058-3063). In addition, several other members of this collagenase gene family have been described recently, including a second type of stromelysin (stromelysin-2), a 92 kilodalton form of type IV collagenase, and Putative Uterine

5 Metalloproteinase (PUMP)-1, a low molecular weight uterine collagenase (Wilhelm, S.M., Collier, I.E., Marmer, B.L., Eisen, A.Z., Grant, G.A., and Goldberg, G.I., 1989, J. Biol. Chem. 264, 17213-17221; Woessner, J. F. and Talpin, C. J., 1988, J. Biol. Chem. 263, 16918-16925).

10 The matrix metalloproteinases are thought to play an important role in disease processes characterized by the inappropriate destruction of the extracellular matrix. The diseases include inflammatory processes such as rheumatoid arthritis and other autoimmune disorders, tumor cell invasion

15 and metastasis formation, local sequelae of myocardial anoxia, and corneal ulceration (Okada et al., 1986, J. Biol. Chem. 261, 14245-14255; Harris et al., 1984, Collagen Relat. 4, 493-512; Werb et al., 1977, New Engl. J. Med. 296, 1017-1023; Liotta et al., 1980, Nature (London) 284, 67-68;

20 Kalebic et al., 1983, Science 221, 281-283). Many tissues contain natural inhibitors of the matrix metalloproteinases. In some cases, this inhibitory activity is derived from the antiproteases in plasma, particularly α_2 -macroglobulin and β_1 -anticollagenase. α_2 -Macroglobulin is a high molecular weight

25 (725,000 Da) inhibitor present in serum. It is thought to account for 95% of the collagenolytic inhibitory activity present in serum. Because of its large size, it is normally unable to pass the vascular permeability barrier. Under conditions of extreme inflammation in which there is

30 increased capillary permeability, α_2 -macroglobulin may enter the tissue compartments and play a role in the regulation of matrix metalloproteinases. The mechanism of inhibition of the matrix metalloproteinases by α_2 -macroglobulin has not been directly studied. However, it is thought to be similar to

35 the mechanism whereby α_2 -macroglobulin causes inhibition of

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other proteases. Hence, the mechanism is not believed to be unique for the matrix metalloproteinases.

β_1 -Anticollagenase is approximately 40,000 daltons in size. It accounts for approximately 5% of the 5 metalloproteinase inhibiting activity of serum. This inhibitor is thought to pass the vascular permeability barrier and be widely distributed in the tissue compartments. β_1 -Anticollagenase may be related to another group of natural 10 inhibitors of the metalloproteinases referred to as TIMPs, tissue inhibitors of metalloproteinases. The down-regulation of metalloproteinase collagenolysis and proteolysis may occur through TIMPs.

The prototype TIMP, TIMP-1, is a glycoprotein with an apparent molecular size of 28.5 kDa which forms a complex of 15 1:1 stoichiometry with activated interstitial collagenase, stromelysin, and the 92 kDa type IV collagenase (Welgus and Stricklin, 1983, J. Biol. Chem. **253**, 12259-12264; Welgus et al., 1985a, Collagen Rel. Res. **5**, 167-179; Wilhelm et al., 1989, J. Biol. Chem. **264**, 17213-17221; European patent 20 189,784). The gene coding for TIMP-1 has been cloned, sequenced and mapped to the X-chromosome (Carmichael et al., 1986, Proc. Natl. Sci. USA **83**, 2407-2411; Docherty et al., 1985, Nature (London) **318**, 66-69; Mullins et al., 1988, Genomics **3**, 187-194; Mahtani et al., 1988, Genomics **2**, 294-25 301). The secreted protein has 184 amino acids and six intramolecular disulfide bonds. Reduction and alkylation of TIMP-1 abolishes all inhibitory activity. The same cells which produce interstitial collagenase are capable of synthesizing and secreting TIMP-1 (Welgus et al., 1985b, J. Clin. Invest. **76**, 219-224; Herron et al., 1986, J. Biol. Chem. **261**, 2814-2818). Thus, the net collagenolytic activity for these cell types is the result of the balance between activated enzyme levels and TIMP-1 levels. Studies have shown an inverse correlation between TIMP-1 levels and the 30 invasive potential of murine and human tumor cells. Down-modulation of TIMP-1 mRNA levels by use of TIMP-1 antisense RNA resulted in conversion of previously nontumorigenic,

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noninvasive Swiss 3T3 cells to tumorigenic cells with invasive properties in vitro and metastatic potential in vivo (Khokha et al., 1989, *Science* 243, 947-950).

Another class of biologically active collagenase 5 inhibitors is composed of low molecular weight (>10,000 daltons) cationic proteins isolated from cartilage, aorta and teeth, but which are poorly characterized.

Recently several new members of the matrix metalloproteinase family have been identified, with various 10 substrate specificities. These include stromelysin (homologue of rat transin), type IV collagenase (70 kDa gelatinase) and a 92 kDa gelatinase. While also identified in normal cell types, the over-expression of these enzymes 15 has been linked to malignant conversion and the metastatic phenotype in a number of systems. Thus, there is a need to understand the molecular basis of the regulation of these metalloproteinases and to find inhibitors which can be exploited for diagnostic and therapeutic purposes.

SUMMARY OF THE INVENTION

20 It is the object of this invention to provide means of purifying natural inhibitors of metalloproteinases.

It is a further object of this invention to provide matrix metalloproteinase inhibitors and derivatives thereof. The inhibitors may be obtained from natural sources, may be 25 produced by synthetic means such as the Merrifield peptide synthesis process, or by genetically engineered organisms or cell lines. The inhibitors of the invention may be used to treat disease conditions which result from activity of matrix metalloproteinases. Furthermore, since metalloproteinase 30 activity is essential to the implantation of the zygote, these inhibitors are useful as contraceptives.

The present invention relates to novel metalloproteinase inhibitors distinct from previous inhibitors mentioned above. Described herein are the 35 isolation and sequencing of a novel protein, now designated TIMP-2 (initially called CSC-21K, so named for its amino terminal amino acid sequence and apparent molecular weight on

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gel electrophoresis). CSC-21K forms a 1:1 complex with type IV procollagenase and type IV collagenase and inhibits activated type IV collagenase. Binding of CSC-21K to activated type IV collagenase results in inhibition of its 5 collagenolytic activity. This inhibitor can be isolated using affinity chromatography on purified matrix metalloproteinase attached to a solid phase. Amino acid sequence analysis of CSC-21K reveals significant homology to TIMP-1 indicating that CSC-21K is the first novel additional 10 member of the family of TIMP-like proteins.

Thus, a preferred embodiment is a protein of approximately 21,600 daltons which binds to matrix metalloproteinases and can be isolated using affinity chromatography on solid phase purified metalloproteinases. 15 The amino acid sequence of this isolated protein shows that it is a new gene product not previously discovered and has areas of sequence homology with the known natural tissue inhibitor of metalloproteinases (TIMP-1). The protein of the preferred embodiment of this inhibitor is characterized by 20 the amino acid sequence shown in Figure 5, below.

DESCRIPTION OF THE FIGURES

Figure 1. Anion exchange chromatography of the complex of collagenase type IV and inhibitor isolated from human melanoma cell (A2058) conditioned media and eluted from 25 gelatin affinity chromatography. 15 µg of gelatin-affinity purified material were applied to the anion exchange resin. The column was eluted with a linear gradient of NaCl (--). Material from the single major peak eluting at 0.18 M NaCl was rechromatographed on the reverse phase column (insert). 30 Material from peaks A and B were sequenced directly (see Fig. 2B).

Figure 2. A. 15% polyacrylamide-SDS gel electrophoresis of CSC-21K and CSC-21K-type IV collagenase complex. Lane A. 2 µg of CSC-21K (peak A) material 35 following reverse phase HPLC purification. Lane B. 2 µg of CSC-21K-type IV collagenase complex isolated by gelatin-

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Sepharose affinity chromatography. Gel was run at 25 milliamps constant current, using a Laemmli sample buffer system and sample buffer containing β -mercaptoethanol. Samples were heated at 95°C for 2 min prior to 5 electrophoresis. B. Amino terminal amino acid sequence of reverse phase HPLC peaks. The complex obtained following gelatin-affinity and anion exchange chromatography was further purified into components by reverse phase HPLC. The materials obtained in peaks A and B (Fig. 1, insert) were 10 sequenced directly.

Figure 3. CSC-21K inhibition of activated type IV collagenase/gelatinase activity. A. Dose relationship of purified CSC-21 (upper curve with, lower curve without reduction and alkylation) inhibition of purified, p-APMA 15 activated type IV collagenase. CSC-21K is termed TIMP-2 and presented as a mole/mole basis. The substrate is native type IV collagen. B. Dose relationship of purified CSC-21K (termed TIMP-2) inhibition of purified p-APMA activated type IV collagenase. The substrate is gelatin.

20 Figure 4. CSC-21K protein sequence data obtained from the amino terminus and following digestions with endoproteinsases Lys-C, Arg-C and Asp-N. Peptide sequences obtained following digests were aligned by overlaps (underlined regions) as shown. The entire sequence of CSC-25 21K is encompassed by these overlapping peptides. The origin of each of the peptides is identified in the lower half of the figure.

Figure 5. Complete sequence for CSC-21K (TIMP-2) derived from direct amino acid sequencing and homology to 30 human TIMP-1. Computerized homology searches using the BIONET system were applied to the sequence obtained following digestions with endoproteinsases Lys-C, Arg-C and Asp-N. The results of these homology searches are shown.

Figure 6. cDNA sequence and deduced protein sequence 35 of clones pSS15 and pSS18 which encode portions of CSC-21K.

Figure 7. Nucleotide sequence and prediction amino acid sequence of a complete human TIMP-2 cDNA. The cDNA

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insert of clone pSS38 was sequenced in both directions using dideoxy-methodology. The predicted amino acid sequence is shown under the DNA sequence. The putative polyadenylation signal is underlined.

5 Figure 8. Comparison of TIMP-2 deduced amino acid sequence and direct amino acid sequencing of CSC-21K protein (see Figure 5, above). CSC-21K primary structure was determined directly using a Porton Instruments 2020 gas phase protein sequenator and phenyl hydantoin derivative 10 identification on a Beckman System Gold HPLC unit equipped with a 0.46 X 25 cm Beckman ODS column. Comparison shows 96% identity of these sequences. Asterisks identify changes in the sequence identified by DNA sequencing of the complete TIMP-2 cDNA.

15 Figure 9. Homology comparison of TIMP-2 and TIMP-1 at the amino acid (A) and nucleotide levels (B). A. Deduced amino acid sequences of TIMP-2 and TIMP-1 were compared using a Pustell Scoring Matrix. The analysis was performed using a cutoff value of 66% homology, and an 8 amino acid overlap. 20 The line denotes regions in which the homology exceeds the average value of 66% homology between these two protein. B. Comparison of the nucleotide sequences of TIMP-2 and TIMP-1. Analysis was performed using a Pustell Scoring Matrix, with a hash value of 4 and a window of 30. The line indicates 25 regions of identity. The analysis if performed for TIMP-1 vs. TIMP-1 or TIMP-2 vs. TIMP-2 gives a solid line on the diagonal, indicating complete identity. This demonstrates that TIMP-2 is a unique gene product distinct from TIMP-1.

30 Figure 10. Northern blot analysis of TIMP-2 mRNA expression in cultured cell lines. Total cytoplasmic and oligo-dT selected RNA as isolated from cells as described in the text. After transfer to Nytran filter RNA was hybridized with ³²P-labelled probe specific for TIMP-2. The resulting autoradiographs and shown. A). Oligo-dT selected RNA (1 μ g) 35 from A2058 human melanoma cells. B). Total cytoplasmic RNA (5 μ g) from WI-38 human embryonic fibroblasts (lane 1) and HT-1080 human fibrosarcoma cells (lane 2).

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Figure 11. Northern blot analysis of total cytoplasmic RNA isolated from A2058 melanoma cells following 48 hour treatment with either 12-tetradecanoylphorbol 13-acetate (10 ng/mL, lane B) or transforming growth factor β 1 (5ng/mL, lane C). These are compared with basal levels in untreated A2058 cells (lane A). Equal amounts of RNA (5 μ g) were loaded and the ethidium bromide stained gel is shown as a control (insert).

Figure 12. Northern blot analysis of human colorectal tumors and adjacent normal mucosa. RNA (5 μ g) of each sample was electrophoresed and transferred as described in the text. Lanes T₁, T₂ and T₃ contain RNA from the invasive colorectal tumors. Lanes N₁, N₂ and N₃ contain RNA from the corresponding adjacent normal mucosa.

15

DETAILED DESCRIPTION OF THE INVENTION

The present invention relates to an inhibitor of matrix metalloproteinases for which the determination of the complete primary structure shows that this protein is a second member of the TIMP family, TIMP-2, as recently 20 reported (Stetler-Stevenson et al., 1989, J. Biol. Chem. **264**, 17374-17378). TIMP-2 is a 21 kDa protein which selectively forms a complex with the latent proenzyme form of the 72 kDa type IV collagenase (Stetler-Stevenson et al., 1989, J. Biol. Chem. **264**, 17374-17378; Goldberg et al., 1989, Proc. Natl. Acad. Sci. USA **86**, 8207-8211). The secreted protein has 192 amino acid residues and is not glycosylated. TIMP-2 shows an overall 71% homology to TIMP-1 at the amino acid sequence level. The position of the twelve cysteine residues are conserved with respect to those present in TIMP-1, as are 30 three of the four tryptophan residues. TIMP-2 inhibits the type IV collagenolytic activity and the gelatinolytic activity associated with the 72 kDa enzyme. Inhibition studies demonstrated that complete enzyme inhibition occurred at 1:1 molar ratio of TIMP-2 to activated 72 kDa type IV 35 collagenase enzyme (Stetler-Stevenson et al., 1989, J. Biol. Chem. **264**, 17374-17378). Thus unlike TIMP-1, TIMP-2 is

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capable of binding to both the latent and activated forms of type IV collagenase. Cell culture studies using cell lines that produce a variety of collagenase family enzymes, as well as both TIMP-1 and TIMP-2, suggest that TIMP-2 preferentially 5 interacts with the 72 kDa type IV collagenase (Stetler-Stevenson et al., 1989, J. Biol. Chem. **264**, 17374-17378; Goldberg et al., 1989, Proc. Natl. Acad. Sci. USA **86**, 8207-8211). Thus, like interstitial collagenase activity which is the balance of activated enzyme and TIMP-1, the net 10 72 kDa type IV collagenase activity may depend upon the balance between the levels of activated enzyme and TIMP-2.

Analogs of the natural inhibitor of the invention can be made by preparing peptides and proteins having cysteines at the same intervals as the cysteines in the natural 15 inhibitor. Other amino acids may vary from the pattern of the natural inhibitor so long as the cysteine is located at the appropriate intervals. At least two appropriately spaced cysteines must be present in the peptide to ensure inhibitory activity by virtue of a disulfide bridge formation.

20 While the preferred protein contains the sequence of Figure 5, peptides having amino acids identical with the sequence of Figure 5 in at least 50% of the sequence positions are within the scope of the invention having useful inhibitor of metalloproteinase activity provided the 25 cysteines are retained in the desired relative positions.

Peptide fragments derived from the natural CSC-21 molecule were used as immunogens. In the case of synthetic peptide fragments of the protein to be used as immunogens or antigens for antibodies specific for CSC-21K, one skilled in 30 the art understands that a unique amino acid sequence in terms of recognition by an antibody binding site consists of a sequence of from four to six amino acids which is not known to exist in another protein in the environment for which the antibody is to be used (e.g., human biological specimens). 35 Further, a unique nucleotide sequence in this context refers to the nucleotide sequence encoding a unique amino acid sequence as defined above and, therefore, consists of from

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four to six codons (12 to 18 nucleotides) needed to encode four to six amino acids. Antibodies to such unique protein fragments can be used to detect the natural inhibitor in serum, tissue, and other natural sources.

5 Particularly preferred peptides are those having at least 2 cysteines. An amino acid sequence containing the sequence CSCSPVHPQQAFCNA derived from the amino terminal of the molecule and segments containing the amino acid sequences 10 SLNHRYQQGCECKITRCP and MIPCYISSLPECLWTD appear particularly active. The antigenic and functional utility of peptides derived from CSC-21K is not limited to these peptides but can include the whole protein, natural or synthetically derived.

Example 1

Purification of CSC-21K (TIMP-2)

15 Human A2058 melanoma cells were grown to 80% confluence in Dulbecco's modified Eagle's medium with 10% fetal bovine serum. The medium was then replaced with serum-free Dulbecco's modified Eagle's medium, and the culture continued for an additional 24 hours. Approximately 60 L of 20 human melanoma cell (A2058) serum-free conditioned medium was concentrated to 300 mL using an Amicon YM 30 ultrafiltration membrane. This concentrated conditioned medium was applied to two 1.0 x 10 cm gelatin-Sepharose (Sigma Chemical Co.) affinity columns in series, equilibrated with 0.05 M Tris 25 HCl, 0.5 M NaCl, 0.005 M CaCl₂, 0.02% Brij 35, pH 7.6 buffer. The columns were then washed with equilibration buffer before eluting with 10.0% DMSO in equilibration buffer. The eluate was concentrated and exchanged into 0.05 M Tris HCl, 0.15 M NaCl, 0.005 M CaCl₂, 0.02% Brij 35, pH 7.6 using an Amicon YM 30 membrane. The samples were stored at -80°C. Samples for 30 anion exchange chromatography were dialyzed into 0.01 M Tris HCl, pH 7.5, with 20% ethylene glycol. A 15 µg sample was injected into a Dionex AI400 HPLC system equipped with a 0.4 x 5.0 cm Dionex ProPac anion exchange column. This column 35 was eluted with a linear gradient of zero to 0.4 M NaCl. The material under the single major peak was collected and an aliquot was applied to a 0.46 x 10 cm RP300 column (Pierce

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Chemical Co.). This column was eluted as previously described (U. S. patent application serial number 07/317,407, filed March 1, 1989, by Liotta et al.). Alternatively, the complex as obtained from gelatin-Sepharose chromatography and 5 stored at -80°C can be applied directly to the RP300 column system.

CSC-21K was isolated as a complex with human type IV collagenase by gelatin affinity chromatography of human melanoma cell (A2058) conditioned media. Anion exchange 10 chromatography of the material obtained from gelatin-affinity chromatography resulted in a single species eluting at approximately 0.18 M NaCl (Figure 1). Reverse phase HPLC analysis of material eluted from the ion exchange chromatography showed that this material contained two 15 components (Figure 1, insert). The material obtained from the gelatin-affinity chromatography step is thus an intermolecular complex as seen on anion exchange chromatography, and is not a simple copurification of two species on gelatin-affinity chromatography. NaDODSO₄-PAGE of 20 the complex obtained from the gelatin-affinity chromatography also showed two components, Figure 2A. The higher molecular weight material has an M_r of 70,000. It was identified as type IV procollagenase by immunoblotting and amino terminal sequencing (*vide infra*). The lower molecular weight material 25 has an apparent M_r of 18,000 which increased to 21,000 upon reduction. Direct reverse phase HPLC analysis of the complex obtained from gelatin affinity chromatography resulted in the separation of two peaks identical to those in the insert of Figure 1. The material obtained from each of these peaks, 30 designated peak A for the material with the shorter retention time and peak B for the material of longer retention time, was subjected to amino acid analysis and direct amino acid sequencing. Peak A material gave a unique amino-terminal amino acid sequence shown in Figure 2B. This material is 35 referred to as CSC-21K. Peak B material gave an amino-terminal sequence identical to latent type IV collagenase (i.e., type IV procollagenase), Figure 2B.

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Example 2

Enzyme digestions, amino acid sequencing, and
amino acid composition analyses

HPLC purified CSC-21K was reduced and alkylated as 5 described. 15 µg of reduced and alkylated CSC-21K was incubated with 5 µg of endoproteinase Lys-C, 5 µg of endoproteinase Arg-C, or 2 µg of endoproteinase Asp-N in 0.1 M NH₄HCO₃ buffer overnight at 37°C. The digests were then separated by reverse phase HPLC on the RP300 column into 10 component peaks which were collected and sequenced individually. Amino acid sequence analysis was carried out on HPLC-purified fractions on a Porton Instruments 2020 Gas Phase Protein Sequenator using standard program 39. PTH amino acid identification was carried out on a Beckman System 15 Gold HPLC unit equipped with a 0.46 x 25 Beckman ODS column and eluted using a modified sodium acetate/THF/acetonitrile separation method.

Amino acid composition analyses were performed following vapor phase hydrolysis for 18 h using 6N HCl, 0.1% 20 phenol at 120°C. The hydrolysate was derivatized using the PITC method (PicoTag system, Waters) and analyzed in the same HPLC unit as above using a modified triethylamine/ammonium acetate/acetonitrile elution method.

Amino acid composition analyses of the complex eluted 25 from the gelatin-affinity chromatography, and CSC-21K are compared in Table 1. The amino acid composition of CSC-21K is significantly different from other collagenase inhibitors and is distinguished by an unusual Leu/Ile ratio. This feature was used to evaluate the stoichiometry of the complex 30 as isolated by gelatin affinity chromatography. Based on the experimentally determined molar amino acid composition of CSC-21 (7 Leu, 18 Ile; Table 1, which is in agreement with the direct amino acid sequence from overlapping peptides, Figure 4) and the deduced composition of type IV 35 procollagenase (39 Leu/25 Ile), it was calculated that the theoretical Leu/Ile ratio of a 1:1 molar complex would be 46

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Leu/42 Ile or 1.10. This is in excellent agreement with the ratio value of 1.03 that was determined from the amino acid composition analysis of the CSC-21K-type IV procollagenase complex. Thus human melanoma cells, which are known to 5 secrete several metalloproteinases, also secrete a protein, CSC-21K, which specifically binds to the latent form of type IV collagenase and forms a complex with 1:1 molar stoichiometry.

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Table 1. Amino acid compositions of CSC-21K (TIMP-2) and the CSC-21K+Collagenase IV Complex

5	AMINO ACID	RESIDUES		
		RESIDUE	Picamoles ⁽¹⁾	MOLE ⁽²⁾
	ASP/ASN		141	18
	GLU/GLN		166	22
10	SER		107	14
	HIS		32	4
	GLY		222	29
	ARG		56	7
	THR		59	8
15	ALA		108	14
	PRO		93	12
	TYR		60	8
	VAL		86	11
	MET		11	1
20	CYS		33	4
	ILE		137	18
	LEU		57	7
	PHE		50	7
	LYS		127	17
25				107
	TOTAL		1545	201
				1361
30	LEU/ILE RATIO		0.42	0.39
				1.02

⁽¹⁾Data obtained from direct amino acid composition analysis of reduced and alkylated CSC-21K, or enzyme inhibitor complex as isolated by gelatin-Sepharose chromatography, as described above.

⁽²⁾Molar amino acid composition calculated from amino acid composition data assuming 7 phenylalanine residues per mole of CSC-21K.

40 The complete primary structure of human CSC-21K, determined by sequence analysis of overlapping peptides obtained following endoproteinase Lys-C, endoproteinase Arg-C and endoproteinase Asp-N digestions, is shown in Figure 4. The amino acid composition of CSC-21K determined from this
45 sequence data concurs with that obtained by direct analysis

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of purified CSC-21K, Table 1. The molecular weight of CSC-21K calculated from the primary sequence is 21,600 daltons, which is in good agreement with the gel electrophoresis data. Computer searches for homology were performed on the BIONET 5 Protein Data Base (which accesses the NBRF-PIR and SWISS-PROT protein sequence data banks). Computerized homology searches were applied to the entire peptide sequence. This provided the basis for alignment of the CSC-21K structure with that of human TIMP, Figure 5. CSC-21K shares significant homology 10 with TIMP. There is 41.0% amino acid identity and 29% conservative substitutions in a 191 amino acid overlap. The positions of the twelve cysteine residues are conserved and the positions of three of four tryptophan residues are also conserved. The conservation of the relative positions of 15 these residues supports their functional or structural roles in both proteins.

Example 3

Collagenolytic and gelatinolytic inhibition.

Type IV collagenase assays were performed as 20 previously described (U. S. patent application serial number 07/317,407, filed March 1, 1989, by Liotta et al.). Gelatinase assays were performed by adaptation of this method utilizing heat denatured rat skin collagen (NEN/Dupont). The CSC-21-collagenase IV proenzyme complex was activated by a 1 25 h preincubation with 1 mM p-aminophenylmercuric acetate (p-APMA). Subsequently, purified CSC-21K was added prior to the assay of collagenase IV activity.

As isolated following gelatin-affinity chromatography, the complex between CSC-21K and type IV procollagenase 30 possessed no collagenolytic activity. Following activation with the organomercurial compound p-amino-phenylmercuric acetate (p-APMA), the maximum achievable type IV collagenolytic activity obtained was 7.12 μ g type IV collagen degraded/h/ μ g enzyme complex. The maximum gelatinolytic 35 activity obtained following organomercurial activation was 26.4 μ g/h/ μ g of enzyme complex. Reduction of disulfide bonds

destroys the complex formed between TIMP and interstitial collagenase and also destroys the complex between CSC-21K and type IV procollagenase (Figure 3A). The addition of purified native CSC-21K, but not reduced and alkylated CSC-21K, to 5 this p-APMA activated complex resulted in proportional inhibition of both collagenolytic and gelatinolytic activities (Figure 3 A and B). Extrapolation of this data demonstrated that the binding of CSC-21K to the activated enzyme occurs in a stoichiometric manner that is consistent 10 with the 1:1 molar ratio determined for the complex isolated by gelatin-Sepharose chromatography. These results demonstrate that CSC-21K which has not been exposed to organomercurial compounds is capable of binding to and inhibiting activated type IV collagenase suggest, but they 15 also suggest that p-APMA activation of type IV collagenase may be accompanied by the organomercurial-mediated inactivation of CSC-21K.

These data show that while CSC-21K shares scattered homology with TIMP-1, particularly with respect to conserved 20 positions of the cysteine residues, all of the CSC-21K peptides are distinctly different from sequences of the known TIMP-1. Thus, the peptides of the invention are encoded by a gene different from that which encodes TIMP. This demonstrates that CSC-21K is the product of a separate gene.

25 Synthetic peptides were prepared using the sequence from the amino terminal portion of the CSC-21K molecule. These were coupled to bovine serum albumin for use in generating anti-peptide antibodies by standard methods. The antibodies were affinity purified using solid phase peptide- 30 affinity chromatography as previously described (Stetler-Stevenson et al., 1989, *J. Biol. Chem.* 264:1353-1356). These antibodies are reactive on standard western and immunoblots.

The isolated, purified CSC-21K, recombinant CSC-21K, and analogs can be used therapeutically in this diseases 35 characterized by the uncontrolled activity of matrix metalloproteinases. Such diseases include arthritis, diabetes, cancer, ulcers of mucosa and epithelial tissues,

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autoimmune mediated inflammation, lung injury, granulomatous diseases. A particularly useful application may be in the treatment of myocardial infarctions since matrix proteolysis including destruction of the myocardial basement membrane is 5 a harmful process in this affliction. Other therapeutic benefit might also be obtained in diseases with basement membrane destruction such as lupus, autoimmune neural disorders, myocyte destruction such as myodystrophies, myocardial infarct and glomerulopathies. CSC-21K could also 10 be used as a potential birth control agent by preventing embryo/placental attachment or invasion.

Example 4

Cloning of Human CSC-21K.

Human A2058 melanoma cells were grown to confluence 15 was run over an oligo dT column to selectively isolate messenger RNA species. This mRNA preparation was then used to prepare a cDNA library using the LambdaGem-4 vector and standard methodology. 1 μ g of purified mRNA was used to prepare double stranded cDNA using a commercially available 20 cDNA synthesis kit (Amersham). This cDNA was methylated using EcoRI methylase (Promega), linked to EcoRI linkers (Promega), restricted with EcoRI and ligated to EcoRI digested Lambda-GEM-4 (Promega). The ligations were packaged (Gigapack Gold, Stratagene) and the optimal reactions were 25 pooled to give 1.5×10^6 recombinants. 7.5×10^5 recombinants were screened using oligonucleotide 27-40. Oligonucleotide 27-40, a 45-mer, with the sequence: 5'-GAGAAGGAGGTGGACTCTGGCAATGACATCTATGGCAACAAACATC-3', corresponding to the reverse translation of residues 27 30 through 40 of the previously sequenced TIMP-2 protein. Oligonucleotide 27-40 was synthesized on a Biosearch 8700 DNA synthesizer by means of β -cyanoethyl phosphoramidite chemistry, and was labelled using γ -[32 P]-ATP (Amersham) and T4 kinase (Bethesda Research Laboratories). From the total 35 of 750,000 plaques screened, 239 positives were identified. Of these positives, initially eight clones were further

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characterized following *SpeI* digestion of the parent LambdaGem-4 clones, religation of the *SpeI* digests and ampicillin resistance selection of the transformants. These eight clones were cross-screened following southern blot hybridization with four additional synthetic oligonucleotides that were also based on the protein sequence data for CSC-21K. Only two clones reacted positively with all four additional synthetic oligonucleotide probes. These clones are designated pSS15 and pSS18. The larger of these two clones is pSS15 which is a pGEM-1 vector containing a 2.1 Kb insert. This clone contains an internal *Hind*III restriction site located approximately 1.2 Kb from the 5' end of the clone. This insert can be released from the pGEM-1 vector by dual endonuclease restriction with *Eco*RI and *Xba*I.

15 Both clones pSS15 and pSS18 were subcloned into M13 and sequenced using the dideoxy method. The pSS15 clone was subcloned using the two *Hind*III fragments. The partial cDNA sequences obtained and the deduced amino acid sequences are shown in Figure 6. The amino acid sequence obtained is identical, within the limits of experimental error, to that obtained for a portion of CSC-21K shown in Figure 5. These results demonstrate that these clones referred to as pSS15 and pSS18 encode the protein CSC-21K. It is obvious that due to differences in codon preferences between species that 25 clones from other species could encode functional CSC-21K protein but with a different nucleotide sequence. Thus, one base change per codon of the CSC-21K cDNA, resulting in 33% change in the overall nucleotide sequence, may still result in a cDNA which would encode a functional CSC-21K protein.

20 30 Thus, the existence of this clone for human CSC-21K is a reduction to practice of isolating the cDNA encoding this protein from other species.

A deposit of CSC-21K cDNA (pSS15) has been made at the American Type Culture Collection (ATCC), 12301 Parklawn Drive, Rockville, Maryland, 20852, U.S.A., on August 11, 1989 under the accession number 40,644. The deposit shall be viably maintained, replacing if it became nonviable, for a

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period of 30 years from the date of the deposit, or for 5 years from the last date of request for a sample of the deposit, whichever is longer, and made available to the public without restriction in accordance with the provisions 5 of the law. The Commissioner of Patents and Trademarks, upon request, shall have access to the deposit.

In subsequent experiments, two addition clones were isolated and the nucleotide sequence of the cDNA insert in the clone pSS38, the longest, is presented in Figure 7. The 10 insert contains 730 bp, excluding the poly(AT) tail and encodes the mature TIMP-2 protein of 194 amino acids. The 130-nucleotide long 3' untranslated region contains the putative polyadenylation signal 30 bases upstream from the 3' end of the RNA.

15 Comparison of the amino acid sequence of TIMP-2 deduced from the cDNA clone with that determined by direct amino acid sequencing of overlapping endoproteinase derived peptide fragments shows excellent agreement. The original sequence contained only 192 amino acids. The previously 20 unidentified residues correspond to the glycyl residue at positions 92 and the prolyl residue at the carboxyl terminus. Other changes are noted in Figure 8. The homology of TIMP-2 with TIMP-2 at the predicted amino acid sequence level is 37.6% identity and 65.6% overall homology. Pustell Matrix 25 analysis of the homology distribution between these two predicted protein sequences using a cutoff value of 66% and an 8 amino acid overlap, demonstrates that there are two areas in which the homology falls below this average value. TIMP-2 shows a distinct preference for binding to the latent 30 form of the 72 kDa type IV collagenase in the presence of both other latent metalloproteinases and TIMP-1 (Stetler- Stevenson et al., 1989, J. Biol. Chem. **264**, 17374-17378; Goldberg et al., 1989, Proc. Natl. Acad. Sci. USA **86**, 8207-8211). However, both forms of TIMP will inhibit 35 activated type IV collagenase. Thus regions of amino acid sequence that are highly conserved between these proteins, such as those that exceed the overall homology value of 66%,

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may be responsible for the known shared functions of these proteins, inhibition of the activated collagenase family enzymes which are unique for individual TIMP molecules. Thus, the regions of low homology between residue 20 to 45 5 and the carboxyl terminus of TIMP-2, may be responsible for the binding of TIMP-2 to the latent form of the 72 kDa type IV collagenase.

Example 5

Applications of the TIMP-2 cDNA.

10 Comparison of the cDNA sequence of human TIMP-2 with that of human TIMP-1 shows little homology compared to that seen at the amino acid level, Figure 9b. This result implies that these genes diverged early in the evolution of this gene family. The lack of homology at the cDNA level may also 15 explain why TIMP-2 mRNA transcripts are not detected in northern blot analyses using TIMP-1 probes and also why screening cDNA libraries with TIMP-1 probes fails to yield TIMP-2 clones.

20 Northern blot analysis of oligo-dT selected mRNA isolated from various cells have been carried out using the TIMP-2 cDNA. HT-1080 human fibrosarcoma cells, WI-38 human embryonic lung fibroblasts, and A2058 human melanoma cells were grown to 80% confluence in Dulbecco's modified Eagle medium (DMEM, GIBCO). The medium was then replaced with DMEM 25 supplemented with 0.5% ITS* (Collaborative Research Inc.) and 25 µg/mL Gentamycin. The medium was changed after 4 hours and culture continued for 20 hours prior to the addition of 10 ng/mL TPA (Sigma Chemical Co.) or 5 ng/mL TGF-β1 (R & D Systems).

30 Total cytoplasmic RNA was isolated from cell lines as described (Gough 1988, Anal. Biochem. 173, 93-95). mRNA was isolated using the FAST-TRACK mRNA isolation kit (Invitrogen). Tissues mRNA was isolated from frozen tissue fragments. Tissue fragments were obtained from three partial colectomy specimens at the time of surgery, from Dr. Barry 35 Schmuckler, Washington Hospital Center, Washington, D.C. The pathologic diagnosis of all three cases was invasive adenocarcinoma. Tissue samples were also obtained from

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adjacent, uninvolved mucosa. Frozen tissue was pulverized in liquid N₂ using a mortar and pestle. The tissue powder was then dissolved in 4 M guanidine isothiocyanate, 3 M sodium acetate, 0.84% β -mercaptoethanol, pH 6.0. Total cytoplasmic 5 RNA was isolated by pelleting through 5.7 M cesium chloride, 3 M sodium acetate, pH 6.0. Aliquots of RNA were applied to formaldehyde/1% w/v agarose gels and electrophoresed before transfer onto Nytran filters (Schleicher & Schuell). The RNA was UV-crosslinked to the filter and hybridized to the insert 10 from clone pSS38. The pSS38 cDNA probe was labelled with α -[³²P]-dCTP using a random primer labelling kit (Bethesda Research Laboratories).

Northern blot analysis of the A2058 human melanoma cell line revealed two specific mRNA species with approximate 15 sizes of 3.5 and 0.9 kb (Figure 10a). These mRNA species were also detected in RNA isolated from human WI-38 fibroblasts, with very low levels of the 0.9 kb species detectable in equivalent amounts of RNA from HT-1080 fibrosarcoma cells (Figure 10b). The origin of these two 20 specific transcripts remains to be determined, however the size difference is too large to be easily accounted for by differences in 3' polyadenylation. It is possible that alternative 5' untranslated regions could account for the different transcript sizes, as has been demonstrated for 25 insulin-like growth factor II mRNA's.

Treatment of A2058 cells with 12-O-tetradecanoylphorbol 13-acetate (TPA) (10 ng/ml) for 48 hours failed to significantly modulate TIMP-2 transcript levels (Figure 11). This is in contrast to mRNA level for the 72 30 kDa type IV collagenase, which was down regulated in response to TPA. Interstitial collagenase mRNA is rapidly induced following TPA treatment of A2058 melanoma cells and fibroblast cells lines (Chin et al., 1985, J. Biol. Chem. 260, 12367-12376; Werb et al., 1986, J. Cell Biol. 102, 697-702; Frisch et al., 1987, Proc. Natl. Acad. Sci. USA 84, 2600-2604), as is TIMP-1 (Edwards et al., 1985, Mol. Cell. Biol. 5, 3280-3288; Murphy et al., 1985, J. Biol. Chem. 260,

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3079-3083; Welgus et al., 1985b, J. Clin. Invest. **76**, 219-224). Treatment of A2058 melanoma cells with transforming growth factor-beta 1 (TGF- β 1) for 48 hours resulted in a clearly detectable decrease in TIMP-2 mRNA levels (Figure 5 11). The 3.5 and 0.9 kb transcripts showed equal decreases in steady-state level, and there was no indication of differential expression. TGF- β 1 has been demonstrated to increase TIMP-1 mRNA levels in human gingival fibroblasts (Overall et al., 1989, J. Biol. Chem. **264**, 1860-1869). It 10 has been demonstrated previously that TGF- β 1 induced the 72 kDa type IV collagenase mRNA and protein levels, as well as enzyme activation. In the presence of the other growth factors, TGF- β 1 also has a selective reciprocal effect on interstitial collagenase and TIMP-1 expression (Edwards et 15 al., 1987, EMBO J. **6**, 1899-1904). TGF- β 1 selectively represses the induction of interstitial collagenase, but interacts synergistically to super-induce TIMP-1. These data demonstrate that TIMP-1 and TIMP-2 respond differently to TPA treatment and oppositely to TGF- β 1 treatment. Furthermore, 20 TGF- β 1 has a reciprocal effect on TIMP-2 and the 72 kDa type IV collagenase transcript levels in human melanoma cells. Thus it is clear that the transcriptional regulation of TIMP-2 is independent of TIMP-1.

Finally, northern blot analysis of tissue from three 25 primary, human colorectal tumors and adjacent normal mucosa was performed using the pSS38 TIMP-2 probe and is presented in Figure 12. The matched samples showed no detectable change in TIMP-2 mRNA transcript levels between the colorectal tumor samples and the adjacent normal mucosa. 30 Previous studies have shown that indeed human colorectal tumor tissue contains elevated type IV collagenase mRNA transcripts. These data suggest that in the primary tumor cell population the ratio of TIMP-2 to 72 kDa type IV collagenase is altered in favor of the enzyme species by 35 differential transcription. However, due to primary tumor cell heterogeneity and the possible inclusion of normal cell

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populations in the invasive tumor samples, this observation may not accurately reflect the invasive, metastatic cell subpopulations. Examination of metastatic lesions will allow a better understanding of the role of these proteins in tumor 5 cell invasion.

Utility of the CSC-21K cDNA Clones

The isolated human cDNA clones encoding the metalloproteinase inhibitor protein, TIMP-2, have wide utility in diagnostics. Pathologic conditions including 10 neoplasia, inflammatory diseases, cardiovascular disease, central nervous system disorders, diabetes and abnormalities of growth and development may be accompanied or causally related to abnormal levels of metalloproteinase inhibitor protein, the subject of invention. All of these processes 15 may involve abnormal accumulation or loss of extracellular matrix proteins. In particular, many of these disease states exhibit abnormal basement membranes. Since control of basement membrane breakdown may be regulated by inhibitors of metalloproteinase action, the inhibitor protein of the 20 subject invention may play a key role in determining the steady-state levels of basement membranes. The present cDNA clones encoding the inhibitor protein CSC-21K can be used in northern blotting analysis to measure the mRNA levels of the inhibitor in RNA samples isolated from tissue samples or 25 cultured cells, as described in Example 5 (above). In some cases, elevated CSC-21K mRNA levels detected in this fashion may reflect pathologic states leading to increased basement membrane accumulation, such as diabetes mellitus. In other cases, loss of the inhibitor protein may be important, such 30 as in neoplasia and central nervous system disorders involving the basement membrane surrounding nerves. In addition to hybridization of the isolated cDNA clone (whole or in part) with isolated RNA or DNA, in situ hybridization using tissue or cell samples can be readily conducted using 35 methods well known in the art. For this and other purposes, the clone can be labeled with radioactive markers for

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detection by using suitable enzymes and radioactive precursors.

The inhibitor protein CSC-21K is a suppressor of neoplastic invasion, and as such, is a tumor suppressor gene.

5 Therefore, homozygous loss, allelic loss, or mutational inactivation of the gene regulatory region may suppress expression of the inhibitor and favor the development of cancer. All of these genetic defects can be detected using the isolated cDNA clone in standard Southern blotting 10 analysis, with or without prior polymerase chain reaction amplification of sample DNA sequences, with methods and appropriate restriction enzymes well known in the art. Extraction of DNA and measurement of such genetic defects may be useful in the diagnosis of cancer and the detection of 15 individuals with hereditary defects predisposing to the development of cancer.

The isolated cDNA clones may be useful in genetic therapy. Diseases associated with loss or down regulation of expression of the subject inhibitor protein could be 20 treated with the cDNA clone in a suitable expression vector, allowing augmented synthesis of the CSC-21K protein. Transfection of the cDNA clone for CSC-21K, in an expression vector with a suitable promoter, into a cell deficient in CSC-21K production would result in the increased production 25 of CSC-21K and correction of the abnormal phenotype. Alternatively, antisense constructs, using the same expression vector but containing the reverse orientation of the CSC-21K cDNA insert, could be used to suppress the overproduction of metalloproteinase inhibitor protein. This 30 could be useful in disorders of abnormal regulation or inappropriately high production of the CSC-21K inhibitor protein. The system and methods of preparation of such genetic reagents is known in the art, but requires the specific isolated nucleotide sequence of the invention. It 35 is obvious that the cDNA clone of the present invention could be spliced next to a gene encoding any other protein to produce a hybrid protein. This methodology could be used to

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produce a hybrid protein with enhanced inhibitory activity or tumor-seeking behavior.

The cDNA clone, or any other DNA segment of this invention which encodes the amino acid sequence of the TIMP-5 2 protein (according to the universal genetic code) is necessary and highly useful for the recombinant production of inhibitor protein CSC-21K (TIMP-2). Recombinant CSC-21K can be made in any suitable expression system, either prokaryotic or eukaryotic. A significant advantage in the 10 production of the present invention is that the protein is not glycosylated, and does not require post-translational modification for functional activity. Thus, recombinant proteins made in bacterial expression systems can be functionally active directly as obtained from the culture 15 medium. It is obvious that the recombinant inhibitor protein can be linked to suitable carrier proteins, marker proteins or other compounds which stabilize or potentiate its activity. The recombinant protein whole, or in part, can be used as an antigen or treatment agent to inhibit 20 metalloproteinase action.

The cDNA clone of the subject invention encodes a novel metalloproteinase inhibitor CSC-21K. The gene itself is novel and distinct encodes a novel metalloproteinase inhibitor CSC-21K. The gene itself is novel and distinct 25 from all prior art reporting cDNA clones encoding proteinase inhibitors. In fact, hybridization of any or all of the cDNA clones for proteinase inhibitors existing in the prior art, under either stringent conditions or conditions of reduced stringency, fails to detect the gene of the present 30 invention, whole or in part. For that reason, the gene of the present invention has never before been detected. The affinity purification of this protein, and the identification of the novel amino acid sequence of this invention was a totally original approach, not described in the prior art 35 which led to the isolation of the gene of the present invention.

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It is well known in the field of protein chemistry that the functional properties of a protein do not depend on the identity of 100% of the amino acid residues which comprise the protein. Individual amino acid residues can be 5 substituted which have the same charge or hydrophobicity and achieve the same function. Furthermore, other amino acids in a protein molecule which specifically function to determine the protein structure can be substituted with residues which differ in charge or hydrophobicity without diminishing the 10 overall biological activity of the parent protein molecule. More generally, the degree of evolutionary similarity of the amino acid sequences of two structurally and functionally related polypeptides is determined by the method of quantitative analysis defined by the sequence alignment and 15 comparison algorithms described by Pearson and Lipman (Pearson, W. R. & Lipman, D. J., 1988, *Proc. Nat. Acad. Sci. U. S. A.* 85:2444-48). This quantitative comparison contemplates not only precise homology of amino acid sequences, but also substitutions of one residue for another 20 which are known to occur frequently in families of evolutionarily related proteins sharing a conserved function. Thus, in the present case, this invention also relates to an isolated polypeptide which inhibits a matrix metalloproteinase and which has an amino acid sequence that 25 differs in at least one position from the sequence defined in Figure 7 and yet has greater similarity to the amino acid sequence of Figure 7, or to a unique portion thereof, than to the amino acid sequence of any other polypeptide.

When matrix metalloproteinase inhibitors of the 30 invention are used in the treatment of inappropriate angiogenesis, arthritis, tumor growth, invasion and metastasis, and granulomatous inflammatory conditions such as sarcoidosis and other pathological conditions, it is possible to estimate the amount of enzyme produced and the 35 amount of peptide inhibitor required to inhibit greater than 90% of the active enzyme. For use in treating any disease condition, the therapeutic dose of the inhibitory peptide

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falls within an acceptable pharmacologic range of 1-250 mg/kg/da, with a more preferred dosage being 25-100 mg/kg/d. The dosage for a given patient will depend on the amount of enzyme produced in the patient, the condition and size of the 5 patient. The inhibitors may be given as infusions or by any means which provides ready transmission into the circulation. Lyophilized powders may be "snorted". Preparations for buccal or sublingual administration may also be given. For respiratory tract involvement, the peptides may be 10 administered by inhalation. Aerosols are particularly useful for this purpose. For conditions of the eye, the peptides may be administered as eye drops.

The isolated CSC-21 proteins, natural or recombinant, or active peptides derived therefrom can be administered 15 intravenously, orally, intrauterine, by inhalation or topical application. For example, topical application can be prepared using a suitable carrier for treatment of basal cell carcinomas or melanomas of the skin or for the treatment of corneal ulceration.

20 The complete CSC-21 protein or CSC-21 peptides can be produced by purification from natural sources, by synthetic peptide chemistry methods or by recombinant DNA technology. In the latter case, suitable cDNA clones for CSC-21 in a suitable expression vector can be used to produce peptides 25 with CSC-21 activity.

CSC-21 peptides and antibodies to CSC-21 are also useful in diagnosis of diseases characterized by abnormal balances of matrix metalloproteinase and associated inhibitor. Purified CSC-21 may be used by virtue of its 30 ability to bind metalloproteinases as a means to purify and or detect metalloproteinases from any natural source. Suitable immunoassays for CSC-21 could include anti-CSC-21 antibodies, reference CSC-21 antigen and solid or solution phase reactions. Purified CSC-21 or peptide domains of CSC-35 21 can be tagged with suitable enzymatic, fluorescent or radioactive labels by means well known in the art.

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Peptides lacking a cysteine or having only one cysteine were found to be useful in assays to detect metalloproteinase and as means of purifying metalloproteinases and are also a part of the invention.

5 Three such structures were peptides having the amino acid sequences:

DIYGNPIKRIQYEIKQIKKFKGIEKDIEFIYTAPSSAVCGVELDVGGK,
DVGGKKEYLIAGKAEDGKRHITL, and RHITLCDFIVPWDTLSTTQKKSLN.

Peptides of the invention may be used in tests to
10 assay metalloproteinases in animal or human tissues or in body fluids which may have antibodies to the protein. Peptides may also be used to elicit antibodies for use in detecting metalloproteinases.

The amino acids herein are given the usual one letter
15 abbreviations accepted as:

A is Alanine	C is Cysteine
D is Aspartic Acid	E is Glutamic Acid
F is Phenylalanine	H is Histidine
I is Isoleucine	K is Lysine
20 L is Leucine	M is Methionine
N is Asparagine	P is Proline
Q is Glutamine	R is Arginine
S is Serine	T is Threonine
V is Valine	W is Tryptophan
25 X is Tyrosine	Y is Pyroglutamic Acid

* * *

For purposes of completing the background description and present disclosure, each of the published articles, patents and patent applications heretofore
30 identified in this specification are hereby incorporated by reference into the specification.

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WHAT IS CLAIMED IS:

1. An isolated polypeptide of approximately 21 kilodaltons which inhibits a matrix metalloproteinase and which binds to both type IV collagenase and type IV procollagenase in a 1:1 stoichiometric ratio.
2. The polypeptide of claim 1 having the amino acid sequence defined in Figure 7, or a unique portion thereof.
3. An isolated polypeptide that inhibits a matrix metalloproteinase, said polypeptide having a unique amino acid sequence that differs in at least one position from the sequence defined in Figure 7, said unique amino acid sequence having greater similarity to the amino acid sequence of Figure 7, or to a portion thereof, than to the amino acid sequence of any other polypeptide.
4. A method of purifying a protein inhibitor of a metalloproteinase having gelatinolytic or collagenolytic activity comprising the steps of
 - (1) attaching a substrate of said metalloproteinase to a solid phase,
 - (2) exposing a mixture of proteins including said inhibitor and said metalloproteinase or the proenzyme form of said metalloproteinase to said immobilized substrate under conditions such that a complex of said inhibitor and said metalloproteinase or said proenzyme will bind to said substrate, and
 - (3) eluting said complex with suitable means.
5. A method of purifying a protein inhibitor of a metalloproteinase having gelatinolytic or collagenolytic activity comprising the steps of
 - (1) attaching said metalloproteinase to a solid phase,
 - (2) exposing a mixture of proteins containing said inhibitor to said immobilized metalloproteinase under conditions such that said inhibitor forms a complex with said metalloproteinase, and
 - (3) eluting said inhibitor with suitable means.

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6. A composition of matter which comprises an amount of said peptide of claim 1 or of claim 3 sufficient to inhibit a matrix metalloproteinase in a pharmaceutically acceptable carrier.

5 7. The composition of claim 6 which is formulated for inhalation.

8. The composition of claim 6 which is a lyophilized powder.

10 9. The composition of claim 6 which is an inclusion complex adapted for buccal or sublingual administration.

10 10. A peptide having the amino acid sequence defined in Figure 7, or a unique portion thereof.

11. The peptide of claim 10 having an amino acid sequence selected from the group consisting of:

15 CSCSPVHPQQAFCNA; SLNHRYQMGCECKITRCP;
MIPCYISSLSPDECLWMD;
DIYGNPIKRIQYEIKQIKMFKGPEKDIEFIYTAPSSAVCGVSLDVGGK;
DVGGKKEYLIAGKAEGDGKMHITL; and MHITLCDFIVPWDLSTTQKKSLN.

12. A composition of matter which comprises said peptide of claim 10 in an amount sufficient to elicit an immune response together with a pharmaceutically acceptable carrier.

20 13. The composition of matter of claim 10 in which said peptide is covalently linked to an immunogenic carrier protein.

14. An antibody elicited in response to administration of a peptide of claim 10.

15. A composition of matter which is at least one antibody of claim 14 attached to a solid support.

30 16. A kit for assaying for a polypeptide of claim 1 in human tissues or body fluids comprising antibodies according to claim 14 and suitable direct or indirect detection means.

17. A kit for assaying for a matrix metalloproteinase 35 in animal or human tissues or body fluids, said kit containing a polypeptide of claim 1 or a peptide of claim 3 as a means to capture or detect said metalloproteinase by

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virtue of binding affinity between said metalloproteinase and said polypeptide or said peptide.

18. A composition of matter comprising a polypeptide of claim 1 or a peptide of claim 3.

5 19. A composition of matter comprising a polypeptide of claim 1 or a peptide of claim 3 attached to a solid support.

20. A DNA segment that encodes a polypeptide of claim 1 or a peptide of claim 3.

10 21. The DNA segment according to claim 20 having the nucleotide sequence shown in Figure 7 or a unique portion thereof.

22. A recombinant DNA molecule comprising said DNA segment according to claim 20 and a vector.

15 23. A culture of cells transformed with said recombinant DNA molecule according to claim 22.

24. A method of inhibiting a matrix metalloproteinase in a mammal comprising administration of a matrix metalloproteinase-inhibiting amount of a composition of claim

20 6.

25. A method of treating a mammal for a degenerative process arising from activity of a matrix metalloproteinase comprising administration of a matrix metalloproteinase-inhibiting amount of a composition of claim 6.

26. The method of claim 25 wherein the matrix metalloproteinase-inhibiting composition is administered sublingually or buccally.

27. The method of claim 25 wherein the degenerative process occurs in the respiratory tract and the matrix metalloproteinase-inhibiting composition is administered by inhalation.

28. The method of claim 26 wherein the matrix metalloproteinase-inhibiting composition is administered as an intravenous infusion.

35 29. A method of inhibiting a matrix metalloproteinase in a mammal comprising administration of a matrix metalloproteinase-inhibiting amount of a polypeptide of claim

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1 or a peptide of claim 3, wherein said polypeptide or peptide is administered by means of administering said recombinant DNA molecule according to claim 22, whereby said DNA molecule is expressed to produce said peptide.

5 30. The method of claim 29 wherein said recombinant DNA molecule is transfected into cells of said mammal followed by administration of the transfected cells to the patient.

10 31. A method of measuring matrix metalloproteinase inhibitor activity comprising the steps of:

- 1) Releasing RNA or DNA from cells or tissue;
- 2) Exposing said RNA or DNA in the preparation of step 1 to a DNA segment of claim 21 under conditions such that said DNA segment will form hybrid duplexes with said RNA or DNA 15 from said cells or tissue; and
- 3) Measuring the amount of said hybrid duplexes resulting from step 2.

32. The method of claim 31 wherein said DNA segment has been labeled whereby detection of said DNA is 20 facilitated.

33. The method of claim 31 wherein exposing said RNA or DNA is performed directly on a preparation of tissue.

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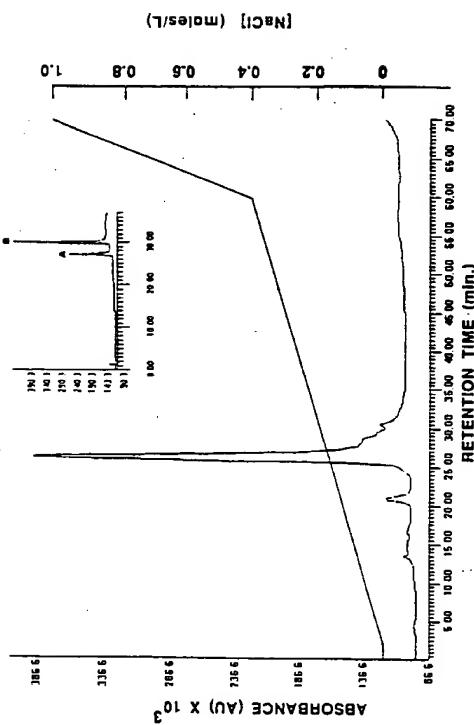


Figure 2.

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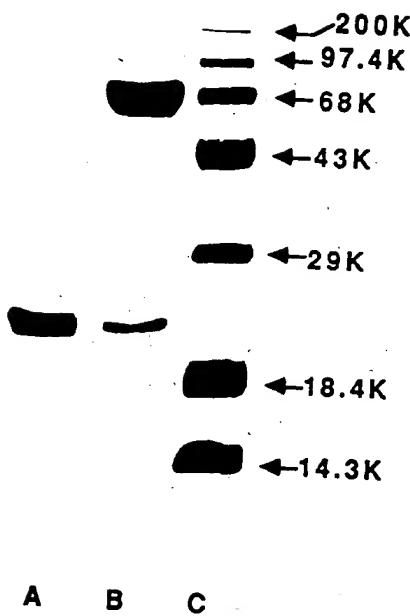
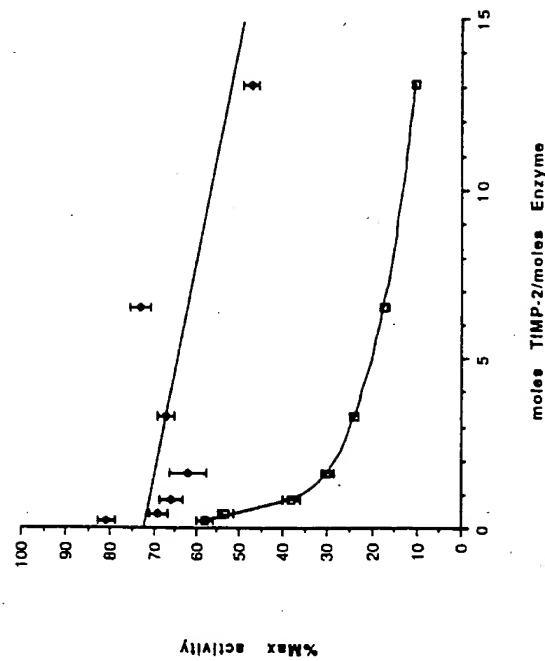


Figure 3.A

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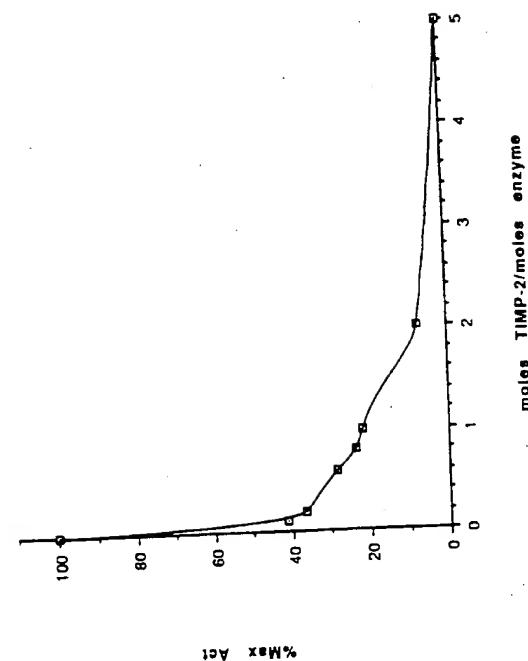


Figure 4.

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SEQUENCE	* OF NEW RESIDUES
*1 CSCSPKHPQQAFNQAVIIRAKVSEKEVOSGMQLYGMPI	40
*2 DIVGMP <u>I</u> ERIQYEV <u>I</u> EQIKFFKGIEK	
*3 GIEKDIEFIV	
*4 DIEFIVTRPSSRUCGU <u>E</u> LGUGK	41
*5 DUGGEKEYLIAKGK <u>E</u>	
*6 EDGKRHI	
*7 DGRKRH <u>I</u> I	18
*8 R <u>H</u> ITLCOFIUPUDTLSTTOKK	
*9 DTLSTTOKK <u>SLM</u>	19
*10 SLMKAVQGCECK	
*11 YQGCECKITR	
*12 ITRCPMIPCYISSPDECLW <u>Q</u> UTEK	36
*13 QHUTEKHMHGK <u>R</u> FFACIKBS	16
*14 RSDOSCAHYVRGARPPK	
*15 GRAPPK <u>E</u> FLD <u>I</u>	
*16 G <u>E</u> FLD <u>I</u> D <u>E</u>	22
	192

SEQUENCE*	DERIVED FROM
1	N-TERMINAL RUN
2,5,6,7,9,13	Asp-N DIGESTION PEPTIDES
3,11,15	Arg-C DIGESTION PEPTIDES
4,8,10,12,14,16	Lys-C DIGESTION PEPTIDES

Figure 5.

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Figure 6.

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A

CCTCCTTGGCTTTCGAAAGCATTTTGGCCAAACTCTTCTCAGGGCTTGATTTCTGAGCTGCGAAATTCTCTGGCTTTCTTAAGCCCTTCCTCC
 ACACCAAGGAGACTCTCTCTCTCTACACCCCTCCGGGAGCCAGCTGCAACAGGCTTCTGAAATGAGATGATGATCAGGGCAAAGGGCTCA 100
 Q Q A F C H A D V V I R A K A V
 CTGAGAGACCAAGTCCACTCTGAAACGACATTATGCCAACCTATCAAGAGGATCCAGTCTGAGATCAGCCAAATAGCTTAAACCCCTGAGA 200
 S H K E V D S G M D I Y G H P I K R I Q Y Z I K Q I K M P F K G P E K
 CGATATAGACTTTCTACACCCCCCTCTCTGGCAACTCTGCTGCTGGGCTTCGGCTGGACCTTGGAGAAAGAACGAAATATCTCTGGAC 300
 D I X F I Y T A P S S A V C G V S L D V G G K E Y L I A C K A Z
 CGGACGCCAAGATGCCACATCACCTCTGACTTCATGGCCCTGAGGACACCCCTGACCAACCCAGAACAGACCCCTGAACCCAGCTGACAC 400
 G D G K M H I T L C D F I V P W D T L S T T Q K S L N E R Y Q M
 CCTGGAGCTCCAAAGATCAGCCCTGCCCATGATCCCTGCTACATCTCCCTCCGGACAGCTCCCTCTGGATGAGCTGGCTCACAGAGAACATCAA 500
 G C E C K I T R C P M I P C Y I S S P D E C L W M D W V T Z K H I N
 CGGGACCCAGGCCAAACTCTTGGCTGCAAGAGAGCTGACCCCTCTGCTGGTACACCCCGCCGGCCCGCCACCCAGACTTCTGGACATC 600
 G B Q A K F F A C I K R S D G S C A W Y R G A A P P P K Q E F L D I
 GAGGACCCATAACGACCCCTCAGACCCCTGCTGAGGAGCTGCAAAAGACCTCCAGGGTTTGGAGGCTCTGAGCTCTGACATCCCTTCTGGAAACAG 700
 E D P
 CATGATAAAACACTCATCCATGGCTCCAAATTATATGAA
 841

441

B.

832

Figure 6. Sequences for cDNA clones pSS15 (A.) and pSS18 (B.).

Figure 7.

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Figure 8.

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CSCSPVHPQQ AFCNADVIR AKA VSEKEVD SGNDIYGNPI KRIQYEIKQI CSC 21K
CSCSPVHPQQ AFCNADVIR AKA VSEKEVD SGNDIYGNPI KRIQYEIKQI TIMP 2

KKFKGIEKDI EFiYTAPSSA VCGVELDVGG KKEYLIAGKA E DGKRHITL CSC 21K
* * * * * KMFKGPEKDI EFiYTAPSSA VCGVSLDVGG KKEYLIAGKA EGDGKMHITL TIMP 2

CDFIVPWDTL STTQKKSLNH RYQQGCECKI TRCPMIPCYI SSPDECLWTD CSC 21K
CDFIVPWDTL STTQKKSLNH RYQMGCECKI TRCPMIPCYI SSPDECLWMD TIMP 2

WVTEKNINGH QAKFFACIKR SDGSCAWYRG APPKQEFID IED CSC 21K
WVTEKNINGH QAKFFACIKR SDGSCAWYRG APPKQEFID IEDP TIMP 2

Figure 9.

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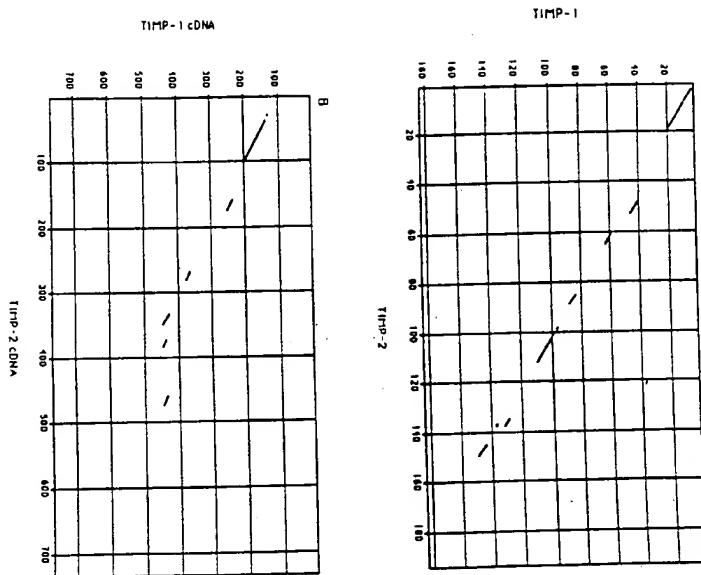


Figure 10.

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A.

B. 1 2

28S>



18S>



Figure 11.

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A B C

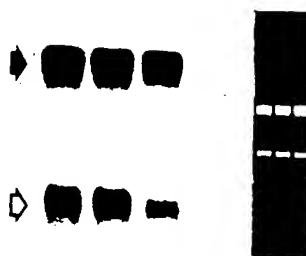


Figure 12.

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 $T_1 N_1 T_2 N_2 T_3 N_3$ 

INTERNATIONAL SEARCH REPORT

International Application No PCT/US90/01526

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all):		
According to International Patent Classification (IPC) or to both National Classification and IPC		
IPC (5): C07H 15/12. C12N 1/22, 15/15 A61K 37/02, 39/00 A01N 37/18		
U.S.CI.: 536/27; 435/252.3, 320; 530/320, 324, 325, 350, 417; 514/2;		
II. FIELDS SEARCHED CONT. 424/85.8		
Minimum Documentation Searched *		
Classification System	Classification Symbols	
U.S.	536/27; 435/252.3, 320 530/324, 325, 350, 417 514/2 424/85.8	
Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched *		
III. DOCUMENTS CONSIDERED TO BE RELEVANT **		
Category *	Citation of Document, if with indication, where appropriate, of the relevant passages **	Relevant to Claim No. **
X	The Journal of Biological Chemistry 263 No. (3) pages 439-1443 (1988) Coulombe et al. "In Vitro Synthesis of the Active Tissue Inhibitor of Metalloproteinases Encoded by a Complementary DNA from Virus-infected Murine Fibroblasts"	1-24
Y		1-24
* Special categories of cited documents: **		
"A" document defining the general state of the art which is not considered to be of particular relevance		
"E" earlier document but published on or after the international filing date		
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)		
"O" document referring to an oral disclosure, use, exhibition or other means		
"P" document published prior to the international filing date but later than the priority date claimed		
"T" later document published after the international filing date but prior to date and not in conflict with the application but cited to understand the principle or theory underlying the invention		
"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step		
"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art		
"Z" document member of the same patent family		
IV. CERTIFICATION		
Date of the Actual Completion of the International Search *		Date of Mailing of this International Search Report *
03 July 1990		30 JUL 1990
International Searching Authority *		Signature of Authorized Officer **
ISA/US		Shelly J. Guest

FURTHER INFORMATION CONTINUED FROM THE SECOND SHEET

V. OBSERVATIONS WHERE CERTAIN CLAIMS WERE FOUND UNSEARCHABLE¹

This international search report has not been established in respect of certain claims under Article 17(2) (a) for the following reasons:

1. Claim numbers _____, because they relate to subject matter not required to be searched by this Authority, namely:

2. Claim numbers _____, because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out¹, specifically:

3. Claim numbers _____, because they are dependent claims not drafted in accordance with the second and third sentences of PCT Rule 8.4(a).

VI. OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING²

This International Searching Authority found multiple inventions in this international application as follows:

I. That species comprising a protein and peptides, method of purification, compositions, an antibody, and a kit classified in Class 530, subclass 326, 324, 325, 350, 417; class 514, subclass 2; class 424, subclass 85.8 (claims 1-19, 24)

see attachment

1. As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims of the international application.

2. As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims of the international application for which fees were paid, specifically claims:

20-23

3. No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claim numbers:

4. As all searchable claims could be searched without effort justifying an additional fee, the International Searching Authority did not invite payment of any additional fee.

Remark on Protest

- The additional search fees were accompanied by applicant's protest.
- No protest accompanied the payment of additional search fees.